LEMON QUINCE- lemon quince spray
Uriel Pharmacy Inc.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Lemon Quince Special

Directions: ONLY FOR USE AS A NASAL SPRAY.

Ages 12 and older: 2-3 sprays. Ages 2-11: 1-2 sprays. Under age 2: Consult a doctor.

Active Ingredients: 100 gm contains: 24 gm Citrus medica e succus 2X; Cydonia e fruct. 2X, Berberis e fruct 3X. Quartz 20X

Inactive Ingredients: Distilled water, Aloe vera juice, Sodium chloride, Grapefruit seed extract

Use: Temporary relief of nasal congestion and allergy symptoms.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use. Do not use if safety seals on box are broken or missing.

SHAKE WELL BEFORE USE.
REFRIGERATE AFTER OPENING.
USE WITHIN 30 DAYS OF OPENING.

Questions? Call 866.642.2858 Uriel, East Troy, WI 53120 shopuriel.com Lot:

LEMON QUINCE
lemon quince spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-6087
Route of Administration	NASAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LEMON JUICE (UNII: AGN709ANTJ) (LEMON JUICE - UNII:AGN709ANTJ)	LEMON JUICE	2 [hp_X] in 1 mL
QUINCE (UNII: 12MCS4H09N) (QUINCE - UNII:12MCS4H09N)	QUINCE	2 [hp_X] in 1 mL
BERBERIS VULGARIS ROOT BARK (UNII: 1TH8Q20J0U) (BERBERIS VULGARIS ROOT BARK - UNII:1TH8Q20J0U)	BERBERIS VULGARIS ROOT BARK	3 [hp_X] in 1 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	20 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CITRUS PARADISI SEED (UNII: 12F08874Y7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-6087-1	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/01/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2009	

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-6087)

Revised: 4/2024 Uriel Pharmacy Inc.